PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter I of the Patent Cooperation Treaty)

(PCT Rule 44bis)

Applicant's or agent's file reference 355492-2970	FOR FURTHER ACTION	See item 4 below		
International application No. PCT/US2004/007017	International filing date (day/month/year) 08 March 2004 (08.03.2004)	Priority date (day/month/year) 07 March 2003 (07.03.2003)]		
International Patent Classification (IPC) or national classification and IPC ⁷ A61L 31/18, A61P 9/00				
Applicant MICRO THERAPEUTICS, INC.				

1.	This international preliminary re International Searching Authori	eport on patentability (C by under Rule 44 bis.1(a	Chapter I) is issued by the International Bureau on behalf of the a).		
2.	2. This REPORT consists of a total of 9 sheets, including this cover sheet. In the attached sheets, any reference to the written opinion of the International Searching Authority should be read as a reference to the international preliminary report on patentability (Chapter I) instead.				
3.	This report contains indications	relating to the following	g items:		
	Box No. I	Basis of the report			
	Box No. II	Priority			
	Box No. III	Non-establishment of applicability	of opinion with regard to novelty, inventive step and industrial		
	Box No. IV	Lack of unity of inve	ention		
	Box No. V	Reasoned statement applicability; citatio	under Article 35(2) with regard to novelty, inventive step or industrial ns and explanations supporting such statement		
	Box No. VI	Certain documents cited			
	Box No. VII	Certain defects in the international application			
	Box No. VIII	Box No. VIII Certain observations on the international application			
4. The International Bureau will communicate this report to designated Offices in accordance with Rules 44bis.3(c) and 93bis.1 but not, except where the applicant makes an express request under Article 23(2), before the expiration of 30 months from the priority date (Rule 44bis.2).					
			Date of important of this manual		
_	Date of issuance of this report 09 September 2005 (09.09.2005)				
The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland		ombettes	Authorized officer Simin Baharlou		
	Facsimile No. +41 22 740 14 35 Telephone No. +41 22 338 71 30				
Form PCT/IB/373 (January 2004)					

PATENT COOPERATION TREATY

RECEIVED 12 AUG 2004

From the		
INTERNATIONAL	SEARCHING	AUTHORITY

WIPO PCT

10:				PCT	
	see form PCT/ISA/220		WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORIT (PCT Rule 43 <i>bis</i> .1)		
			Date of mailing	e form PCT/ISA/210 (second sheet)	
A	oplicant's or agent's file reference		ļ	·	
<u> </u>	ee form PCT/ISA/220		FOR FURTHER A See paragraph 2 belo	ACTION w	
Ini	ernational application No. CT/US2004/007017	International filing date (o		Priority date (day/month/year)	
	·	08.03.2004	·	07.03.2003	
Af	ernational Patent Classification (IPC) or b 51L31/18, A61P9/00	oth national classification a	and iPC	L	
 					
	plicant CRO THERAPEUTICS, INC.				
1.	This opinion contains indication	ns relating to the follo	wing items:		
	Box No. I Basis of the opin		-		
	Box No. II Priority				
	Box No. III Non-establishme	ent of opinion with regar	d to novelty inventive	step and industrial applicability.	
	En Land of Chilly Of 1	IIIA G III(O)}			
	Box No. V Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement			ovelty, inventive step or industrial	
	Box No. VI Certain documer		supporting such state	ment	
	Box No. VII Certain defects in	n the international applic	cation		
	Box No. VIII Certain observations on the international application				
2.	FURTHER ACTION		,,,		
	If a demand for international preilm written opinion of the International the applicant chooses an Authority International Bureau under Rule 66 will not be so considered.	., ., ., ., ., ., ., ., ., ., ., ., ., .	TOWNSHIP LAND, MO	WRVEL THIS GOOD BOLDS IN THE	
	If this opinion is, as provided above submit to the IPEA a written reply to months from the date of mailing of I whichever expires later.	, considered to be a writ	tten opinion of the IPI	=A the applicant in implication	
	For further options, see Form PCT/				
3.	For further details, see notes to For				

Name and mailing address of the ISA:

Authorized Officer

European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465

Allnutt, S

Telephone No. +49 89 2399-7817



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Box No. I Basis of the oninion
 With regard to the language, this opinion has been established on the basis of the international application in the language in which it was field, unless otherwise indicated under this item.
☐ This opinion has been established on the basis of a translation from the original language into the following language , which is the language of a translation furnished for the purposes of international search (under Rules 12.3 and 23.1(b)).
With regard to any nucleotide and/or amino acid sequence disclosed in the international application and necessary to the claimed invention, this opinion has been established on the basis of:
a. type of material:
□ a sequence listing
☐ table(s) related to the sequence listing
b. format of material:
☐ in written format
☐ in computer readable form
c. time of filing/furnishing:
☐ contained in the international application as filed.
filed together with the international application in computer readable form.
furnished subsequently to this Authority for the purposes of search.
In addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed or furnished, the required statements that the information in the subsequent or additional appropriate, were furnished.
4. Additional comments:

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	Box	k No. II	Priority
1.	×	The fol	lowing document has not been furnished:
		\boxtimes	copy of the earlier application whose priority has been claimed (Rule 43bis.1 and 66.7(a)).
			translation of the earlier application whose priority has been claimed (Rule 43bis.1 and 66.7(b)).
			quently it has not been possible to consider the validity of the priority claim. This opinion has neless been established on the assumption that the relevant date is the claimed priority date.
2.		has be	oinion has been established as if no priority had been claimed due to the fact that the priority claim en found invalid (Rules 43 <i>bis</i> .1 and 64.1). Thus for the purposes of this opinion, the international atteindicated above is considered to be the relevant date.
3.	Ado	ditional c	bservations, if necessary:

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Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability					
			tion appears to be novel, to involve an inventive step (to be non nave not been examined in respect of:		
	the entire international application,				
Ø	claims Nos. 17-21				
bed	ause:				
Ø	the said international application, or the said claims Nos. 17-21 (industrial applicability) relate to the following subject matter which does not require an international preliminary examination (specify):				
	see separate sheet				
	the description, claims or drawings (indicate particular elements below) or said claims Nos. are so unclear that no meaningful opinion could be formed (specify):				
	the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.				
	no international search report has been established for the whole application or for said claims Nos.				
	the nucleotide and/or amino acid sequence listing does not comply with the standard provided for in Annex C of the Administrative Instructions in that:				
	the written form		has not been furnished		
			does not comply with the standard		
	the computer readable form		has not been furnished		
			does not comply with the standard		
	the tables related to the nucleon not comply with the technical re	tide a equire	and/or amino acid sequence listing, if in computer readable form only, do ements provided for in Annex C-bis of the Administrative Instructions.		
	See separate sheet for further	detai	ls ·		

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Box No. V Reasoned statement under Rule 43*bis*.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)

Yes: Claims

No: Claims

1-23

Inventive step (IS)

Yes: Claims

No: Claims

1-23

Industrial applicability (IA)

Yes: Claims

1-16,22,23

No: Claims 17-21

2. Citations and explanations

see separate sheet

Item III

- 1. Claims 17-21 relate to subject-matter considered by this Authority to be covered by the provisions of Rule 67.1(iv) PCT. Consequently, no opinion will be formulated with respect to the industrial applicability of the subject-matter of these claims (Article 34(4)(a)(i) PCT).
- 2. The following documents are referred to in this communication; the numbering will be adhered to in the rest of the procedure:
 - D1: US-A-5 667 767 (EVANS SCOTT ET AL) 16 September 1997 (1997-09-16)
 - D2: US-A-5 695 480 (EVANS SCOTT ET AL) 9 December 1997 (1997-12-09)
 - D3: WO 00/71170 A*(ROTH NOAH M ; GREFF RICHARD J (US); TRAN CHINH N (US); WHALEN THOMAS J) 30 November 2000 (2000-11-30)
 - D4: WO 00/56370 A (DOLMATCH BART; RICCI CHARLIE (US); CRAGG ANDREW H (US); GREFF RICHARD) 28 September 2000 (2000-09-28)
 - D5: US 2002/187102 Å1 (EVANS SCOTT ET AL) 12 December 2002 (2002-12-12)
 - D6: US-B-6 454 7381 (WHALEN II TOM ET AL) 24 September 2002 (2002-09-24)
 - D7: US-A-5 925 683 (PARK SANGSOO) 20 July 1999 (1999-07-20)
 - D8: MOTTU F ET AL. "Iodine-containing cellulose mixed esters as radiopaque polymers for direct embolization of cerebral aneurysms and arteriovenous malformations." January 2002 (2002-01), BIOMATERIALS. JAN 2002, VOL. 23, NR. 1, PAGE(\$) 121 131, XP002291570 ISSN: 0142-9612
 - D9: WRIGHT K C ET AL: "Experimental evaluation of cellulose acetate NF and ethylene-vinyl alcohol copolymer for selective arterial embolization."

 JOURNAL OF VASCULAR AND INTERVENTIONAL RADIOLOGY: JVIR. OCT 1999, vol. 10, no. 9, October 1999 (1999-10), pages 1207-1218, XP000800903 ISSN: 1051-0443

The documents considered in the present processing are consecutively numbered D1-D9; this numbering results from the citations D1-D9 found in the Search Report (SR) of the corresponding PCT application. It will be adhered to in the rest of the procedure. The cited passage(s) for each citation will be considered

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unless otherwise specified.

Item V

Novelty

3. The technical features of claims 1-23 are disclosed by documents D1-D8 and therefore lack novelty in terms of Art 33 (2) PCT.

D1 discloses embolizing compositions comprising a polymer, solvent and up to about 40 wt% of a contrast agent. The compositions are said to form a coherent solid material and the particle size of the contrast agent is maintained at about $2\mu m$ to enhance formation of the suspension.

D2 provides embolic compositions with improved visualization using contrast agent particle sizes of less than 10 μ m. The contrast agent is present at up to 40 wt%. D3, D5, D6 also discloses embolizing compositions comprising a polymer, solvent and up to 40 wt% of a contrast agent.

D4 provides example compositions containing 40 wt% contrast agent. The ratio of polymer to contrast agent is 0.0625 and 0.2 for examples E and F respectively. D7 discloses in example 1, a composition comprising 46%wt of a contrast agent, ultravist 370. The polymer composition obstructed blood flow when injected into the ret mirabile.

D8 appears to provide iodine containing polymer compositions with sufficient radiopacity for use as embolization materials. Iodine was present in amounts of 41.4 and 44.7 wt%.

4. D9 is a general review of known products Embolyx C and E containing 30 %wt tantalum and was not considered to anticipate the subject matter of the claims.

Further Remarks:

Industrial Applicability (Art 33(4) PCT).

5. For the assessment of the present claims 17-21 on the question whether they are industrially applicable, no unified criteria exist in the PCT Contracting States. The patentability can also be dependent upon the formulation of the claims. The EPO, for example, does not recognize as industrially applicable the subject-matter of claims to the use of a compound in medical treatment, but may allow, however, claims to a

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known compound for first use in medical treatment and the use of such a compound for the manufacture of a medicament for a new medical treatment.